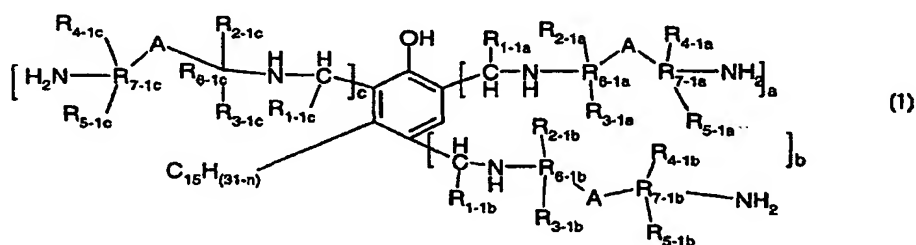




INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁷ : C07C 215/50, C08G 59/40, 59/62	A1	(11) International Publication Number: WO 00/01659 (43) International Publication Date: 13 January 2000 (13.01.00)
(21) International Application Number: PCT/EP99/04323 (22) International Filing Date: 22 June 1999 (22.06.99) (30) Priority Data: 60/091,467 1 July 1998 (01.07.98) US (71) Applicant: CIBA SPECIALTY CHEMICALS HOLDING INC. [CH/CH]; Klybeckstrasse 141, CH-4057 Basel (CH). (72) Inventors: CHENG, Chi-Wen, Frank; 5 Hallmark Drive, New City, NY 10956 (US). BEDNER, David; Prospect Street, Portchester, NY 10573 (US). WANG, Hsing, Tie; 105-09 62 Drive, Forest Hills, NY 11375 (US).		(81) Designated States: AU, BR, CN, JP, KR, European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE). Published With international search report.

(54) Title: PHENALKYLAMINE DERIVATIVES, THEIR USE AS CURING AGENTS IN EPOXY RESIN COMPOSITIONS AND CURABLE EPOXY RESIN COMPOSITIONS CONTAINING THEM



(57) Abstract

The present invention relates to a compound according to formula (1), wherein n is 0, 2, 4 or 6 and a, b and c are, independently of one another, 1 or 0. R_{1-1a}, R_{1-1b} and R_{1-1c} are hydrogen, a hydrocarbyl containing 1 to 10 carbon atoms which are alkyl, aryl, alkylene, arylalkyl or alkylaryl or a hydrocarbyl containing 1 to 10 carbon atoms and at least one heteroatom which can be oxygen, sulfur or nitrogen. R_{2-1a}, R_{3-1a}, R_{4-1a}, R_{5-1a}, R_{2-1b}, R_{3-1b}, R_{4-1b}, R_{5-1b}, R_{2-1c}, R_{3-1c}, R_{4-1c} and R_{5-1c} hydrogen or C₁-C₄alkyl. R_{6-1a}, R_{7-1a}, R_{6-1b}, R_{7-1b}, R_{6-1c} and R_{7-1c} are C₁-C₄alkyl. A is an aromatic or alicyclic ring having 5 to 7 carbon atoms. The present invention further relates to a Mannich base reaction product obtained by combining an extract from cashew nutshell liquid with at least one aromatic or alicyclic polyamine and at least one aldehyde compound, epoxy resin compositions and curable formulations containing the same, and methods for using such compositions.

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece	ML	Mali	TR	Turkey
BG	Bulgaria	HU	Hungary	MN	Mongolia	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MR	Mauritania	UA	Ukraine
BR	Brazil	IL	Israel	MW	Malawi	UG	Uganda
BY	Belarus	IS	Iceland	MX	Mexico	US	United States of America
CA	Canada	IT	Italy	NE	Niger	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NL	Netherlands	VN	Viet Nam
CG	Congo	KE	Kenya	NO	Norway	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NZ	New Zealand	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	PL	Poland		
CM	Cameroon	KR	Republic of Korea	PT	Portugal		
CN	China	KZ	Kazakhstan	RO	Romania		
CU	Cuba	LC	Saint Lucia	RU	Russian Federation		
CZ	Czech Republic	LI	Liechtenstein	SD	Sudan		
DE	Germany	LK	Sri Lanka	SE	Sweden		
DK	Denmark	LR	Liberia	SG	Singapore		
EE	Estonia						

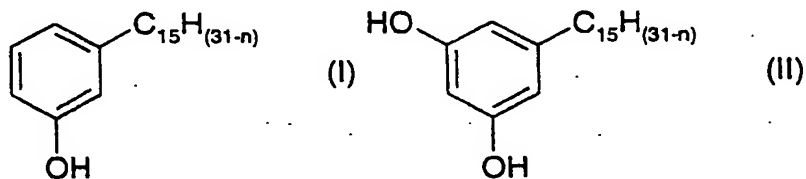
PHENALKYLAMINE DERIVATIVES, THEIR USE AS CURING AGENTS IN EPOXY RESIN COMPOSITIONS AND CURABLE EPOXY RESIN COMPOSITIONS CONTAINING THEM

The present invention relates to a novel class of phenalkamines and epoxy resin compositions containing the same wherein the cured coatings resulting therefrom have reduced Gardner color index. In particular, the novel phenalkamines are prepared via a Mannich base reaction sequence by reacting a selected phenol compound that is substituted with a carbonyl group containing compound and at least one selected aromatic or alicyclic polyamine. The resulting phenalkamine can be used as a curing agent in epoxy resin compositions for coatings, finishes and varnishes.

Mannich base reactions are well-known. Mannich base compounds are products based on the reaction of an aldehyde, generally formaldehyde, a phenolic compound and an amine. Various forms of phenolic compounds, amines and aldehydes have been proposed. Mannich base products are known to be suitable for curing epoxy resins.

Phenalkamine curing agents are a class of Mannich bases obtained by reacting a cardanol-containing extract derived from cashew nutshell liquid, an aldehyde compound, such as formaldehyde, and an amine. Commercially available phenalkamines use ethylenediamine and diethyltriamine as the amine. Phenalkamines are good epoxy resin hardeners for room temperature or low temperature curing applications. Phenalkamines, however suffer from the disadvantage that the cured epoxy articles are very dark in color (Gardner Color Index greater than 14), blushing and low mechanical properties. Due to the dark coloring of the cured coatings, epoxy resin compositions containing phenalkamine curing agents have been limited to use as primers and generally cannot be used as top coat material.

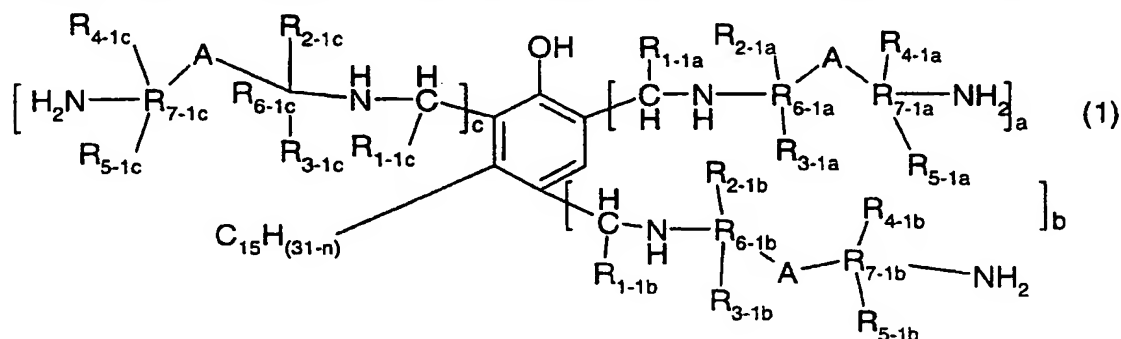
The extract from cashew nutshell liquid primarily contains a mixture of cardanol (I), cardol (II),



where n depends upon whether the side chain is saturated or unsaturated (n=0 for saturated, n=2 for monoene, n=4 for diene, and n=6 for triene), and related compounds of varying degrees of saturation. Numerous methods have been developed to purify the

extract from cashew nutshell liquid with the stated goal of obtaining an isolated solution of cardanol. Typically, a purified extract solution will contain primarily cardanol with a minor amount of cardol.

The present invention relates to a novel phenalkamine compound according to formula (1)



wherein n is 0, 2, 4 or 6,

a, b and c are, independently of one another, 1 or 0, R_{1-1a}, R_{1-1b} and R_{1-1c} are, independently of one another, hydrogen, a hydrocarbyl containing 1 to 10 carbon atoms which are alkyl, aryl, alkylene, arylalkyl or alkylaryl or a hydrocarbyl containing 1 to 10 carbon atoms and at least one heteroatom which can be oxygen, sulfur or nitrogen;

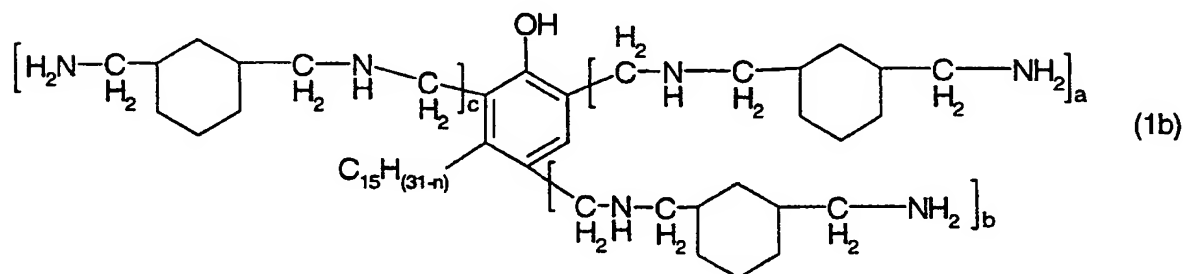
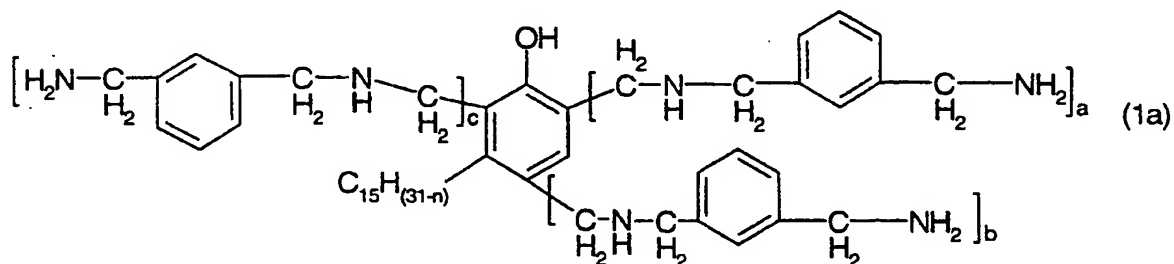
R_{2-1a}, R_{3-1a}, R_{4-1a}, R_{5-1a}, R_{2-1b}, R_{3-1b}, R_{4-1b}, R_{5-1b}, R_{2-1c}, R_{3-1c}, R_{4-1c} and R_{5-1c} are independently of one another, hydrogen or C₁-C₄alkyl, R_{6-1a}, R_{7-1a}, R_{6-1b}, R_{7-1b}, R_{6-1c} and R_{7-1c} are, independently of one another C₁-C₄alkyl; and A is an aromatic or alicyclic ring having 5 to 7 carbon atoms. Preferably, R_{1-1a}, R_{1-1b} and R_{1-1c} are independently of one another, methyl, and a, b and c, independently of one another are 1 or 0. More preferably, R_{1-1a} is hydrogen, a is 1 and b and c are 0.

R_{2-1a}, R_{3-1a}, R_{4-1a}, R_{5-1a}, R_{2-1b}, R_{3-1b}, R_{4-1b}, R_{5-1b}, R_{2-1c}, R_{3-1c}, R_{4-1c} and R_{5-1c} are preferably, independently of one another, hydrogen, methyl or ethyl, and a, b and c, independently of one another are 1 or 0. More preferably, R_{2-1a}, R_{3-1a}, R_{4-1a}, and R_{5-1a} are each hydrogen, a is 1 and b and c are 0. Most preferably, R_{2-1a}, R_{3-1a}, R_{4-1a}, and R_{5-1a} are each hydrogen, a is 1 and b and c are 0.

R_{6-1a}, R_{7-1a}, R_{6-1b}, R_{7-1b}, R_{6-1c} and R_{7-1c} are preferably, independently of one another, methyl or ethyl and a, b and c, independently of one another, are 1 or 0. More preferably, R_{6-1a} and R_{7-1a} are methyl or ethyl, a is 1 and b and c are 0.

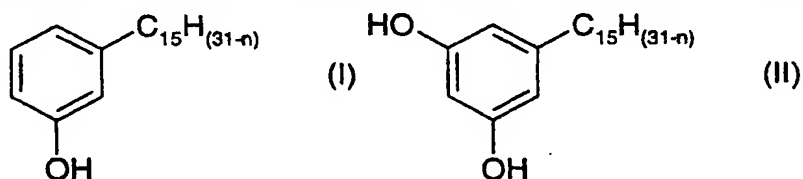
A is preferably an aromatic or alicyclic ring having 6 carbon atoms. More preferably, A is an aromatic or alicyclic ring having 6 carbon atoms.

The present invention is a compound ideally represented by formulae (1a) or (1b):

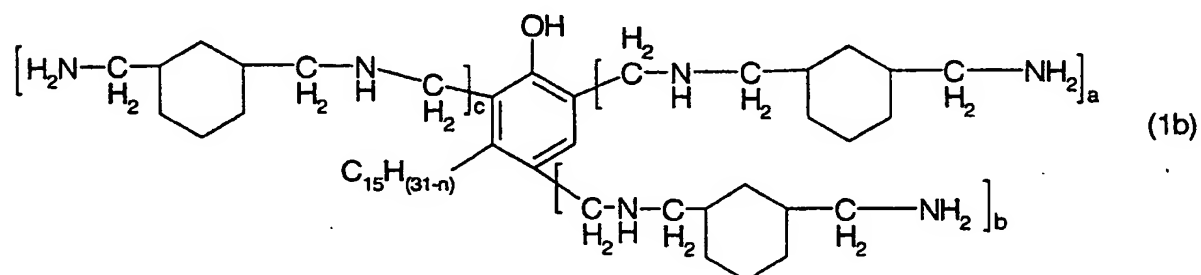
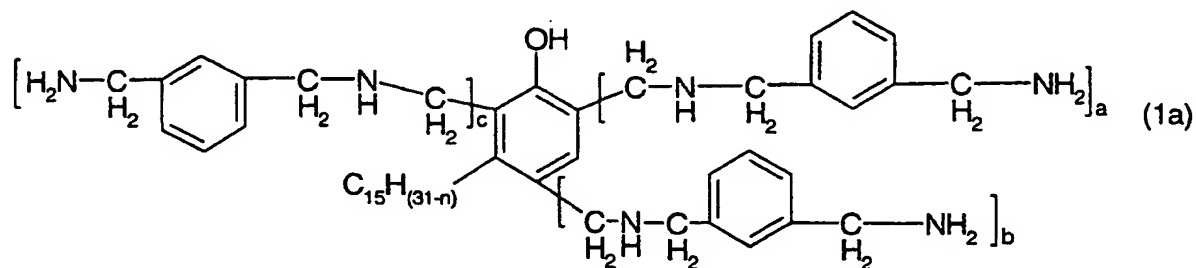


wherein n, a, b and c have been defined above.

The present invention further relates to a Mannich base reaction product obtained by combining an extract from cashew nutshell liquid with at least one aromatic or alicyclic polyamine and at least one aldehyde compound. The extract cashew nutshell liquid preferably contains a major portion of cardanol (I) and a minor amount of cardol (II)



where n is 0, 2, 4 or 6. The extract cashew nutshell liquid preferably contains cardanol (I) and of cardol (II) in a weight ratio of about 90:10 to about 98:2. The Mannich base reaction product is preferably obtained by combining the extract and the at least one aldehyde compound with a polyamine selected from xylenediamine, 1,3-bis(aminomethyl)cyclohexane, and mixtures thereof. The Mannich base reaction product preferably comprises at least one compound ideally represented by structural formulae (1a) or (1b):



wherein n has been defined above, a , b and c , are, independently of one another, 1 or 0, and an average value for the sum of $(a+b+c)$ is about 1.2 for all of the compounds corresponding to formulae (1a) and/or (1b) in said Mannich base reaction product mixture.

The present invention further relates to an epoxy resin composition including a) a novel phenalkamine compound described above and b) an epoxy resin having on average more than one glycidyl group per molecule. Alternatively, the present invention relates to an epoxy resin composition including a) a Mannich base reaction product described above and b) an epoxy resin having on average more than one glycidyl group per molecule.

The present invention further relates to a curable formulation including a) an epoxy resin composition containing the novel phenalkamine compound and an epoxy resin having on average more than one glycidyl group per molecule and b) a pigment. In the alternative, the present invention relates to a curable formulation including an epoxy resin composition containing a Mannich base reaction product and an epoxy resin having on average more than one glycidyl group per molecule and b) a pigment.

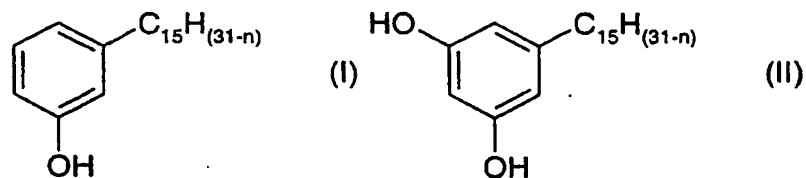
The present invention further relates to a method for coating a surface having reduced yellowness index using a phenalkamine-containing composition that comprises applying a curable formulation described above to the surface. In the alternative, the present invention relates to a method for coating a surface having reduced yellowness index using a Mannich

base reaction product-containing composition that comprises applying a curable formulation described above.

The novel Mannich base curing agents of the present invention can be prepared by reacting an extract derived from cashew nutshell liquid containing at least one phenolic compound with an aldehyde compound and at least one selected aromatic- or alicyclic-polyamine, or mixtures thereof. The mole ratio of polyamine to phenolic compound is within the range of about 1:1 to about 10:1, more preferably from about 1:1 to about 4:1. The mole ratio of the polyamine to aldehyde compound is within the range of about 1:1 to about 5:1, preferably about 1:1 to about 2:1. On an equivalents basis, the ratio of aldehyde and amine should be more than or equal to one mole of amine per equivalent of the phenolic compound.

The polyamine and extract containing the phenolic compound are placed in a suitable vessel, and mixed thoroughly. The aldehyde compound is then added, either continuously over a period of time, or incrementally. The reaction is nearly spontaneous and fairly exothermic. Provisions for temperature control are necessary. After completion of aldehyde compound addition, water that is formed during synthesis is removed by distillation. Chemical reaction during synthesis is believed to involve electrophilic addition of the aldehyde to the phenolic compound to form an alkanolated phenol intermediate. Further condensation with the amine and elimination of water yields the Mannich base reaction product.

The liquid cashew nutshell extract solution, after heat treatment, contains a mixture of cardanol (I), cardol (II), and related low molecular weight compounds, wherein the mixture comprises a major portion of cardanol (I) and a minor amount, of cardol (II)

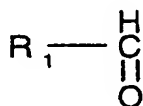


where n depends upon whether the side chain is saturated or unsaturated (n=0 for saturated, n=2 for monoene, n=4 for diene, and n=6 for triene), and related compounds of varying degrees of saturation. Preferably, the ratio of cardanol (I) relative to cardol (II) is about 90:10. The extract solution can be obtained by heat treatment of the cashew nutshell

liquid or oil. Numerous methods have been developed to purify the extract from cashew nutshell liquid with the stated goal of obtaining an isolated solution of cardanol.

The extract solution can be purified by vacuum distillation to further reduce the amount of cardol in the extract solution. The crude cashew nutshell liquid is available in countries, such as Brazil and India. Palmer International Inc., which is located in Worcester, Pennsylvania, is a major importer of crude cashew nutshell liquid. The crude cashew nutshell liquid can be vacuum distilled using commercial distillation facilities to obtain a purified extract solution described above. Palmer is capable of producing such a purified extract solution containing at least about 85% by weight of a mixture containing a major portion of cardanol and a minor amount of cardol.

The aldehyde compound can be any compound containing the group (C=O) which occurs in aldehydes. These compounds can be characterized by the structural formula:

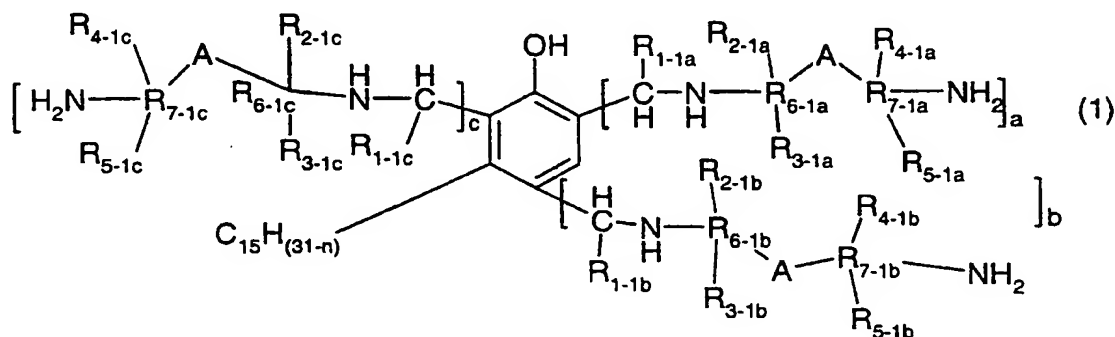


in which R₁ is hydrogen or a hydrocarbyl containing 1 to 10 carbon atoms which can be alkyl, aryl, alkylene, arylalkyl or alkylaryl. The hydrocarbyl groups can also contain 1 to 10 carbon atoms and at least one heteroatom which can be oxygen, sulfur or nitrogen. Typical aldehyde compounds are formaldehyde, butyraldehyde, heptaldehyde, hexadecyde, acetaldehyde, propionaldehyde, paraformaldehyde, benzaldehyde, salicylaldehyde and 2-ethylhexanal. Formaldehyde (in solution or as paraformaldehyde) is particularly preferred. These compounds are known in the art and are readily available from commercial sources or are easily made using known methods.

The polyamines used herein to produce the inventive compounds are selected from aromatic or alicyclic polyamines or mixtures thereof. The polyamine is preferably m-xylenediamine or liquid mixtures thereof with p-xylenediamine, and/or, 1,3- or 1,4-bis(aminoalkyl)cyclohexane or mixtures thereof. The aminoalkyl group is preferably an aminomethyl, aminoethyl, aminopropyl or aminobutyl, wherein the alkyl group is either a straight chain or branched. More preferably, the aminoalkyl group is aminomethyl or aminoethyl. The polyamine is particularly preferably m-xylenediamine alone, or in combination with p-xylenediamine, or

1,3-bis(aminomethyl)cyclohexane, which are commercially available from Mitsubishi Gas and Chemical, Tokyo, Japan.

The resulting novel Mannich base reaction product corresponds to a compound according to formula (1)



wherein

n is 0, 2, 4 or 6,

a, b and c are, independently of one another, 1 or 0,

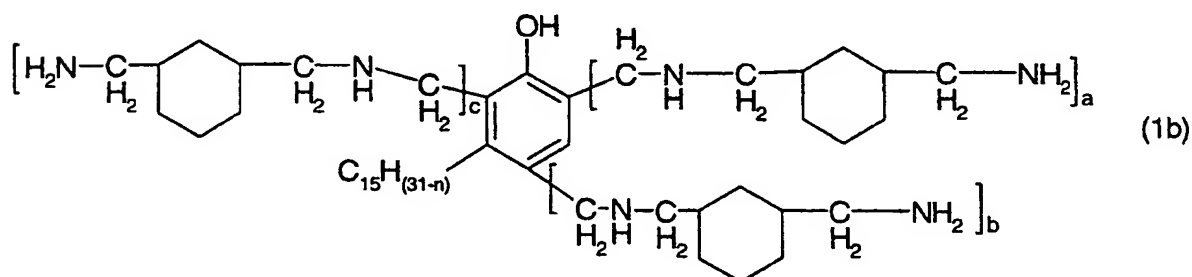
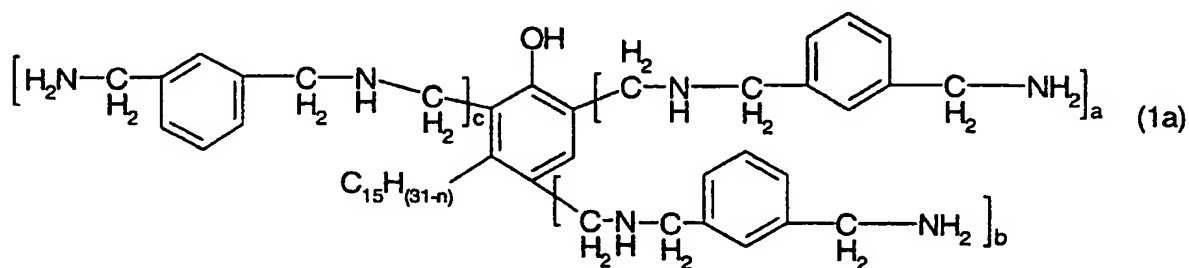
R_{1-1a}, R_{1-1b} and R_{1-1c} are, independently of one another, hydrogen, a hydrocarbonyl containing 1 to 10 carbon atoms which are alkyl, aryl, alkylene, arylalkyl or alkylaryl or a hydrocarbonyl containing 1 to 10 carbon atoms and at least one heteroatom which can be oxygen, sulfur or nitrogen;

R_{2-1a}, R_{3-1a}, R_{4-1a}, R_{5-1a}, R_{2-1b}, R_{3-1b}, R_{4-1b}, R_{5-1b}, R_{2-1c}, R_{3-1c}, R_{4-1c} and R_{5-1c} are independently of one another, hydrogen or C₁-C₄alkyl,

R_{6-1a}, R_{7-1a}, R_{6-1b}, R_{7-1b}, R_{6-1c} and R_{7-1c} are, independently of one another C₁-C₄alkyl; and

A is an aromatic or alicyclic ring having 5 to 7 carbon atoms.

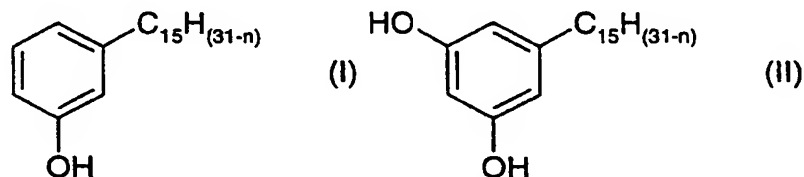
A particularly preferred compound according to the present invention is derived from a reaction mixture containing formaldehyde as the aldehyde compound and xylenediamine or 1,3(bisaminomethyl)cyclohexane as the polyamine. Hence, a particularly preferred compound can be represented by the following idealized structural formulae (1a) and (1b):



wherein n, a, b and c have been defined above.

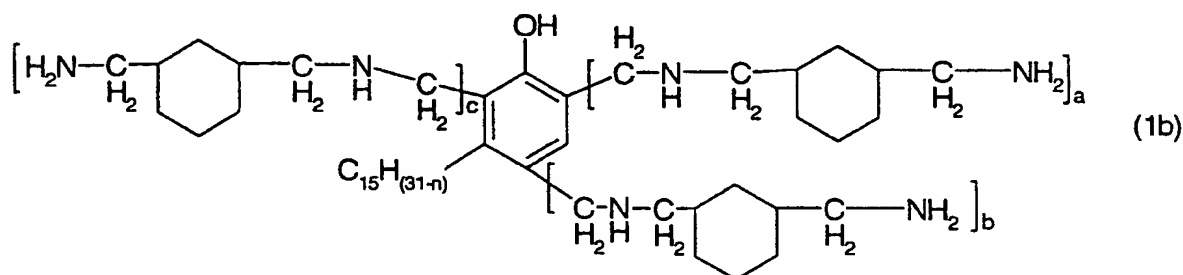
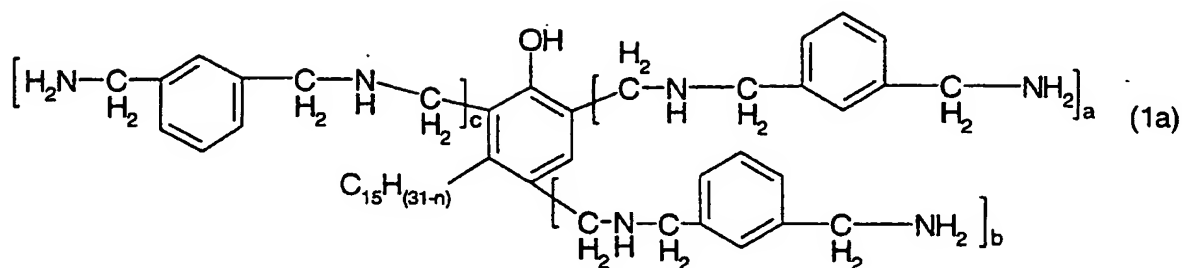
The term "hydrocarbyl" as used herein encompasses any hydrocarbon radical, whether saturated, unsaturated, linear or cyclic subject to any other restrictions apparent from the text.

The novel compounds described above are obtained via a Mannich base reaction sequence using an extract of cashew nutshell liquid or oil. The cashew nutshell liquid can be obtained in varying degrees of purity. Accordingly, the novel compound will preferably be produced from a reaction mixture containing an extract having a major portion of cardanol (I) and a minor amount of cardol (II)



where n is 0, 2, 4 or 6.

A particularly preferred reaction mixture will contain said extract and at least one aldehyde compound, preferably formaldehyde, with a polyamine selected from xylenediamine, 1,3-bis(aminomethyl)cyclohexane, and mixtures thereof. The product mix of said reaction mixture will contain at least the following compounds ideally represented by structural formulae (1a) and (1b):



wherein n has been defined above and average value for the sum of $(a+b+c)$ is about 1.2 based upon an average of all of the compounds corresponding to formulae (1a) and/or (1b) in said resulting product mix. The average value for the sum of $(a+b+c)$ is determined from the molar ratio of the alkylated phenol to aldehyde to amine, which is most preferably 1:1.5:1.5.

The resulting Mannich base reaction products can be used as a curing agent for epoxy resin compositions. The epoxy resin can be any epoxy resin which can be cured by a Mannich base curing agent. Generally, the epoxy resin can be any curable epoxy resin having at least one glycidyl group per molecule. The epoxy resin can be saturated or unsaturated, linear or branched, aliphatic, cycloaliphatic, aromatic or heterocyclic, and may bear substituents which do not materially interfere with the curing reaction. Such substituents can include bromine. The epoxy resin can be monomeric or polymeric, liquid or solid, but is preferably liquid at room temperature. Suitable epoxy resins include glycidyl ethers prepared by reacting epichlorohydrin with a compound containing at least one, preferably two or more, hydroxyl groups carried out under alkaline reaction conditions. Examples of epoxy resins suitable for use in the invention include polyglycidyl ethers of polyhydric phenols, epoxy novalacs or similar glycidylated polyphenolic resins, polyglycidyl ethers of alcohols, glycols or polyglycols, and polyglycidyl esters of polycarboxylic acids, and mixtures thereof.

The preferred epoxy resin is a resin based on a polyglycidyl ether of a polyhydric phenol. Polyglycidyl ethers of polyhydric phenols can be produced, for example, by reacting an epichlorohydrin with a polyhydric phenol in the presence of an alkali. Examples of suitable polyhydric phenols include: 2,2-bis(4-hydroxyphenyl)propane (bisphenol A); 2,2-bis(4-hydroxy-3-tert-butylphenyl)propane; 1,1-bis(4-hydroxyphenyl)propane; 1,1-bis(4-hydroxyphenyl)ethane; 1,1-bis(4-hydroxyphenyl) isobutane; bis(2-hydroxy-1-naphthyl)methane; 1,5-dihydroxynaphthalene; 1,1-bis(4-hydroxyphenyl)isobutane; 1,1-bis(4-hydroxy-3-alkylphenyl) ethane and the like. Suitable polyhydric phenols can be obtained from the reaction of phenol with aldehydes, such as formaldehyde (bisphenol F). The preferred epoxy resin can be a mixture of the above resins. Particularly preferred epoxy resins are based on bisphenol A, such as GY 6010, available from Ciba Specialty Chemicals Corporation, Brewster, New York.

The preferred epoxy resins can also be blended with a glycidyl ether of an aliphatic or aromatic alcohol, glycol or polyglycol, or glycidylester of a monocarboxylic acid. Examples include butyl glycidyl ether, phenyl glycidyl ether, cresyl glycidyl ether, 1,4-butanediol, diglycidyl ether, the glycidyl ester of neodecanoic acid, and the like. Such glycidyl ethers and esters can be blended with the preferred epoxy resins in concentrations of from about 1 to about 50% by weight in order to affect properties, such as wetting, viscosity, flexibility, and adhesion.

The preferred epoxy resin, which can be a mixture of the resins described above, can be combined with a curing system containing the novel Mannich base reaction product described above. The curing system can be combined with the epoxy resin at a temperature below the curing temperature. The preferred epoxy resin or curing system can be diluted with minor amounts of aliphatic or aromatic hydrocarbons, alcohols, or glycol ethers to facilitate handling and mixing requirements.

The epoxy resin composition can optionally further include other additives, such as flow control additives, antifoam agents, anti-sag agents, pigments, reinforcing agents, fillers, elastomers, stabilizers, extenders, plasticizers or flame retardants.

The resulting curable composition can be cured at a temperature within the range of from about -40°C, preferably from about -10°C, to about 150°C, for a sufficient time to fully cure

the epoxy resin. For standard ambient cure applications, the composition is preferably cured at a temperature of from about -5°C to about 40°C.

The epoxy resin composition described herein can be used as an adhesive, coating, flooring, casting or encapsulants, to name a few applications. The epoxy resin composition has particularly good applicability as a top coating, especially when combined with a pigment. The epoxy resin composition using the novel Mannich base reaction products described above can advantageously be combined with a pigment to produce a paint formulation having a reduced Gardner color index relative to currently available paint formulations containing phenalkamines. The resulting formulations can be applied in conventional manner by spray, roller or brush, for instance.

The novel Mannich base reaction products described above and particularly preferred methods for preparing and using said products are discussed more fully below in the following non-limiting examples.

Example 1 - Synthesis of Phenalkamine with m-xylenediamine

Charge 600 grams (2.0 moles) of cardanol (vacuum distilled cashew nutshell liquid to have a minimum purity of 85% cardanol and cardol, ratio of cardanol to cardol 92:8) obtained from Palmer International and 396 grams (2.9 moles) m-xylenediamine obtained from Mitsubishi Gas and Chemical, into a 2-liter, 3-necked round bottom flask equipped with a thermometer, a mechanical agitator, and a Dean-Stark water trap connected to a condenser. Then add 96 grams (3.20 moles of formaldehyde) of 92-94% paraformaldehyde (6-8% by weight water) (molecular weight 30) obtained from Aldrich Chemical Co. over a 20 minute period at a rate that maintains a temperature in the range of 80-100°C. A water bath can be used to maintain the desired temperature, particularly not in excess of 100°C, in the reaction vessel. Over a four-hour period, raise the temperature from 100°C to 120°C to remove and recover 52 grams of water in the water trap. The reaction completion can be monitored by way of an amine number titration, which shows 300-330 mg KOH/g. Upon completion, cool the pot temperature to 80 to 100°C for discharge of 1040 grams of a clear red liquid having an amine value of 315 mg KOH/gram, Gardner color of 16, and a viscosity at 25°C of 3,000 mPa·s (centipoise).

Example 2 - Synthesis of Phenalkamine with 1,3-bis(aminomethyl)cyclohexane

Charge 600 grams (2.0 moles) of cardanol (vacuum distilled cashew nutshell liquid) described above and 406 grams (2.9 moles) 1,3-bis(aminomethyl)cyclohexane obtained from Mitsubishi Gas and Chemical into a 2-liter, 3-necked round bottom flask equipped with a thermometer, a mechanical agitator, and a Dean-Stark water trap connected to a condenser. Then add 105 grams (3.50 moles of formaldehyde) of 92-94% paraformaldehyde (6-8% by weight water) (molecular weight 30) obtained from Aldrich Chemical Co. over a 20 minute period at a rate that maintains a temperature in the range of 80-100°C. A water bath can be used to maintain the desired temperature, particularly not in excess of 100°C, in the reaction vessel. Over a four-hour period, raise the temperature from 100°C to 120°C to remove and recover 56 grams of water in the water trap. The reaction completion can be monitored by way of an amine number titration (290-320 mg KOH/g). Upon completion, cool the pot temperature to 80 to 100°C for discharge of 1055 grams of a clear red liquid having an amine value of 295 mg KOH/gram, Gardner color of 16, and a viscosity at 25°C of 14,000 centipoise.

Example 3 - Blushing Test

The following formulations (amounts are in parts by weight unless stated otherwise) were prepared by mixing the resin and hardener components for about 2 minutes to produce a uniform paste.

Component	Gardner Color	Formulation #3-1	Formulation #3-2	Formulation #3-3	Formulation #3-4
GY 6010	<1	100	100	100	100
HY3440	16	43.3			
HY3441	16		69.6		
Product of Example 1	16			69.6	
Product of Example 2	16				68

GY 6010 is an unmodified liquid epoxy resin based on bisphenol A and epichlorohydrin having a viscosity of 11,000 to 14,000 centipoise at 25°C and an epoxy equivalent weight of about 182-192 grams per equivalent. GY 6010 is commercially available from Ciba Specialty Chemicals Corporation, Brewster, New York.

HY3440 is a phenalkamine based on an extract of liquid cashew nutshell liquid (85% purity with ratio of cardanol to cardol of 92:8) and formaldehyde wherein the polyamine is diethyltriamine.

HY3441 is a phenalkamine based on an extract of liquid cashew nutshell liquid (85% purity with ratio of cardanol to cardol of 92:8) and formaldehyde wherein the polyamine is ethylenediamine.

The resulting formulations were then coated as a 6 mil wet film onto a glass plate. The coated glass plates were cured in a refrigerator at 5°C for 4 to 8 hours. The plates were then removed from the refrigerator and placed in a controlled environment at 25°C overnight with the following results:

Component	Formulation #3-1	Formulation #3-2	Formulation #3-3	Formulation #3-4
Results	5	4	1	0

where on a scale of 0 to 5, 0 means no blush and 5 means worst blush.

Example 4

The following formulations (amounts are in parts by weight unless stated otherwise) were prepared by mixing the resin and hardener components for about 2 minutes to produce a uniform paste.

Component	Gardner Color	Formulation #4-1	Formulation #4-2	Formulation #4-3	Formulation #4-4
GZ9625 W90	<1	100	100	100	100
HY3440	16	31			
HY3441	16		49.8		
Product of Example 1	16			49.8	
Product of Example 2	16				48.6

GZ9625 W90 is an unmodified bisphenol A "Type 1/2" epoxy having a 90% solids content in an n-butanol solution and having an epoxy equivalent weight of about 500 grams per equivalent. GZ9625 W90 is commercially available from Ciba Specialty Chemicals Corporation, Brewster, New York.

The resulting formulations were then coated as a 6 mil wet film onto a glass plate. The coated glass plates were cured in a refrigerator at 5°C for 4 to 8 hours. The plates were

then removed from the refrigerator and placed in a controlled environment at 25°C or 5°C overnight to determine appearance. Tack-free and dry-through times were determined using Gardner drying time recorder. The tack-free time is the point in time when the ball scribe of the Gardner recorder penetrates the film and the groove exposes an underlying metal substrate. The dry-through time is the point in time in which the ball scribe can no longer make a groove on the film as evidenced by the end of the groove. Pencil hardness is determined by ASTM method No. D3363-92a. Cross-hatch adhesion is determined by ASTM method No. D3359-95. Impact is determined by ASTM method No. D2794. The Mandrel bend is determined by ASTM method No. D522-93a. Each of the above ASTM test methods are incorporated herein by reference.

Component	Formulation #4-1	Formulation #4-2	Formulation #4-3	Formulation #4-4
Appearance at 5°C	blush	slight blush	glossy	glossy
Appearance at 25°C	slight blush	glossy	glossy	glossy
Tack-free at 25°C, hr.	2	2.5	2	2
Dry-through at 25°C, hr.	3.5	3.5	3.5	4
Tack-free at 5°C, hr.	5	3	2	2.5
Dry-through at 5°C	19	19	7	11
Pencil Hardness	4H	2H	2H	2H
Cross Hatch adhesion (5 is best)	5	5	5	5
Impact (pass) direct, lb.	24	24	20	10
Impact (pass) reverse, lb.	6	2	0	0
Mandrel bend	pass	fail	fail	fail

Example 5

The following formulations (amounts are in parts by weight unless stated otherwise) were prepared by mixing the resin and hardener components for about 2 minutes to produce a uniform paste.

Component	Gardner Color	Formulation #5-1	Formulation #5-2	Formulation #5-3	Formulation #5-4
GY 6010	<1	100	100	100	100
HY3440	16	43.3			
HY3441	16		69.6		
Product of Example 1	16			69.6	
Product of Example 2	16				68

Component	Formulation #5-1	Formulation #5-2	Formulation #5-3	Formulation #5-4
Appearance at 5°C	blush	slight blush	glossy	glossy
Appearance at 25°C	slight blush	slight blush	glossy	glossy
Tack-free at 25°C, hr.	3.5	3	3	3.5
Dry-through at 25°C, hr.	9.5	8.5	5.5	5
Tack-free at 5°C, hr.	5	4	4	5
Dry-through at 5°C	23	19	14	13
Pencil Hardness	3H	B	3H	2H
Cross Hatch adhesion (5 is best)	5	5	5	5
Impact (pass) direct, lb.	20	26	20	10
Impact (pass) reverse, lb.	0	0	0	0
Mandrel bend	fail	pass	fail	fail

Example 6

The following formulations (amounts are in parts by weight unless stated otherwise) were prepared by mixing the resin, hardener and pigment components for about 2 minutes to produce a uniform paste.

Component	Gardner Color	Formulation #6-1	Formulation #6-2	Formulation #6-3	Formulation #6-4
R28P-40		100	100	100	100
GZ9625 W90	<1	15.5			
HY3440	16	35.8			
HY3441	16		49.8		
Product of Example 1	16			49.8	
Product of Example 2	16				48.6
Pigment/Binder	40/60	40/60	40/60	40/60	

RP28P-40 is a white paint formulation containing 100 parts by weight of GZ9625 W90 and 93.2 parts by weight of titanium dioxide, TI-Pure, R706, available from DuPont, Wilmington, Delaware. The resulting formulations were then coated as a 10 mil wet film onto a cool roll steel panel. The "in the light" panels are left on the laboratory bench and exposed to fluorescent light. The "in the dark" panels are stored in drawers to exclude any light. The coated steel panels were cured overnight at 22°C and monitored for yellowness with the following results:

In the Light	Formulation #6-2	Formulation #6-3	Formulation #6-4
Day 1	23.28	13.16	15.12
Day 2	23.47	13.99	16.58
Day 3	24.64	15.49	17.32
Day 6	25.94	16.43	19.66
Day 17	26.99	18.56	22.85
Day 24	26.75	19.93	23.59
Day 31	26.82	20.53	24.3

In the Dark	Formulation #6-2	Formulation #6-3	Formulation #6-4
Day 1	25.43	12.13	12.26
Day 2	25.31	11.98	12.39
Day 3	26.68	12.99	13.9
Day 6	27.11	13.17	14.73
Day 17	26.61	13.12	16.31
Day 24	27.44	14.07	17.32
Day 31	26.14	14.47	18.53

Example 7

The following formulations (amounts are in parts by weight unless stated otherwise) were prepared by mixing the resin, hardener and pigment components for about 2 minutes to produce a uniform paste.

Component	Formulation #7-1	Formulation #7-2
R28P-20A	100	100
HY3441	49.8	
Product of Example 2		28.3
Pigment/Binder	40/60	40/60

RP28P-20A is a white paint formulation containing 363.6 grams of GY 6010, 400 grams of titanium dioxide (R-706), available from DuPont and 72.7 grams of n-butanol. The resulting formulations were then coated as a 10 mil wet film onto a cool roll steel panel. The coated steel panels were cured overnight at 22°C and monitored for yellowness with the following results:

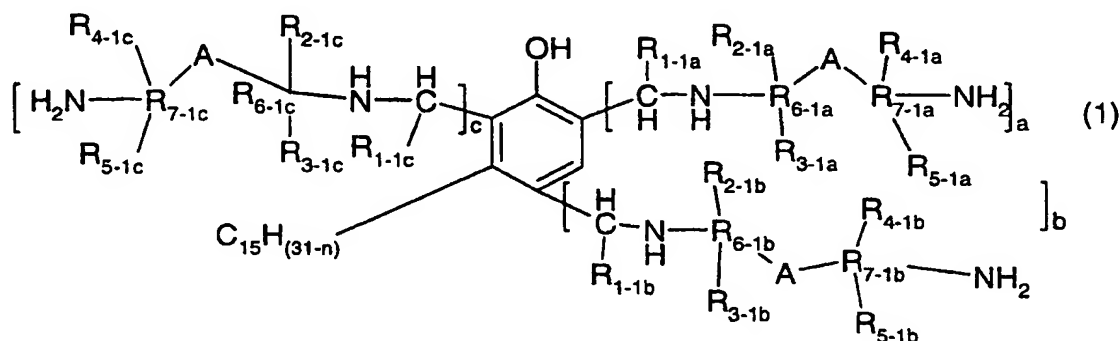
In the Light	Formulation #7-1	Formulation #7-2
Day 1	26.07	18.71
Day 2	29.06	21.57
Day 4	31.03	23.08
Day 8	32.17	23.88
Day 16	33.48	25.35
Day 31	35.31	27.70
Day 65	36.75	29.10
Day 110	38.81	31.06
Day 209	46.54	35.74

In the Dark	Formulation #7-1	Formulation #7-2
Day 1	26.07	18.71
Day 2	26.52	19.64
Day 4	26.77	19.89
Day 8	26.95	19.99
Day 16	27.88	20.49
Day 31	28.98	21.03
Day 65	30.72	21.63
Day 110	31.88	22.18
Day 209	34.53	23.45

Preferred embodiments of the present invention relating to novel phenalkamines, compositions thereof and methods for using the same have been described above. Those skilled in the art having the benefit of the teachings presented in the foregoing will recognize modifications and other embodiments. Therefore, it is understood that the invention is not limited to the specific embodiments disclosed herein, and that modifications and other embodiments are intended to be within the scope of the appended claims.

Claims

1. A compound according to formula (1)



wherein

n is 0, 2, 4 or 6,

a, b and c are, independently of one another, 1 or 0,

R_{1-1a}, R_{1-1b} and R_{1-1c} are, independently of one another, hydrogen, a hydrocarbyl containing 1 to 10 carbon atoms which are alkyl, aryl, alkylene, arylalkyl or alkylaryl or a hydrocarbyl containing 1 to 10 carbon atoms and at least one heteroatom which can be oxygen, sulfur or nitrogen;

R_{2-1a}, R_{3-1a}, R_{4-1a}, R_{5-1a}, R_{2-1b}, R_{3-1b}, R_{4-1b}, R_{5-1b}, R_{2-1c}, R_{3-1c}, R_{4-1c} and R_{5-1c} are independently of one another, hydrogen or C₁-C₄alkyl,

R_{6-1a}, R_{7-1a}, R_{6-1b}, R_{7-1b}, R_{6-1c} and R_{7-1c} are, independently of one another C₁-C₄alkyl; and

A is an aromatic or alicyclic ring having 5 to 7 carbon atoms.

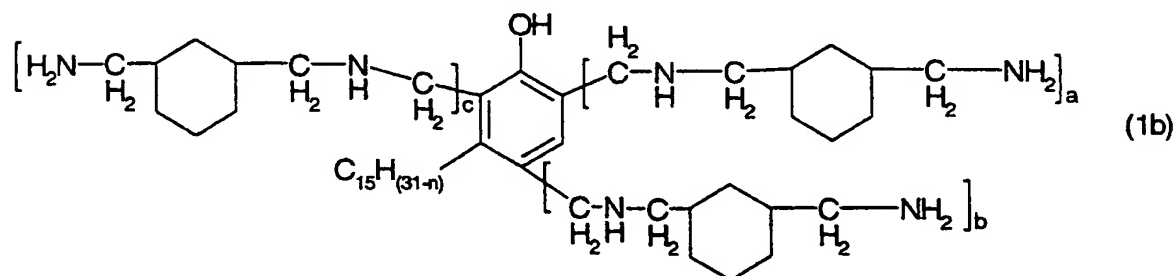
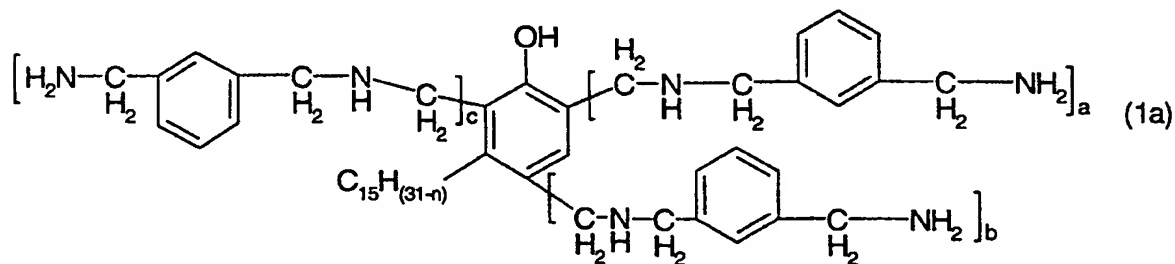
2. A compound according to claim 1 wherein R_{1-1a}, R_{1-1b} and R_{1-1c} are independently of one another, methyl, and a, b and c, independently of one another are 1 or 0.

3. A compound according to claim 2 wherein R_{1-1a} is hydrogen, a is 1 and b and c are 0.

4. A compound according to claim 1 wherein R_{2-1a}, R_{3-1a}, R_{4-1a}, R_{5-1a}, R_{2-1b}, R_{3-1b}, R_{4-1b}, R_{5-1b}, R_{2-1c}, R_{3-1c}, R_{4-1c} and R_{5-1c} are, independently of one another, hydrogen, methyl or ethyl, and a, b and c, independently of one another are 1 or 0.

5. A compound according to claim 4 wherein R_{2-1a}, R_{3-1a}, R_{4-1a}, and R_{5-1a} are each hydrogen, a is 1 and b and c are 0.

6. A compound according to claim 3 wherein R_{2-1a} , R_{3-1a} , R_{4-1a} , and R_{5-1a} are each hydrogen, a is 1 and b and c are 0.
7. A compound according to claim 1 wherein R_{6-1a} , R_{7-1a} , R_{6-1b} , R_{7-1b} , R_{6-1c} and R_{7-1c} are, independently of one another, methyl or ethyl and a, b and c, independently of one another, are 1 or 0.
8. A compound according to claim 6 wherein R_{6-1a} and R_{7-1a} are methyl or ethyl, a is 1 and b and c are 0.
9. A compound according to claim 1 wherein A is an aromatic or alicyclic ring having 6 carbon atoms.
10. A compound according to claim 8 wherein A is an aromatic or alicyclic ring having 6 carbon atoms.
11. A compound according to claim 1 represented by formulae (1a) or (1b):

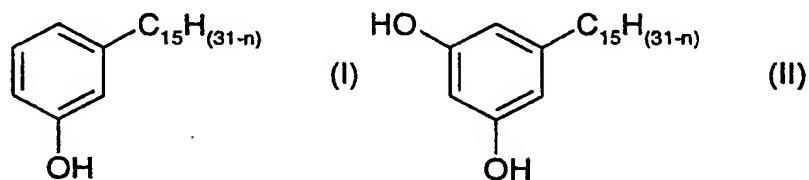


w

herein n, a, b and c have been defined above.

12. A Mannich base reaction product obtained by combining an extract from cashew nutshell liquid with at least one aromatic or alicyclic polyamine and at least one aldehyde compound.

13. A Mannich base reaction product according to claim 12 wherein the extract cashew nutshell liquid contains a major portion of cardanol (I) and a minor amount of cardol (II)

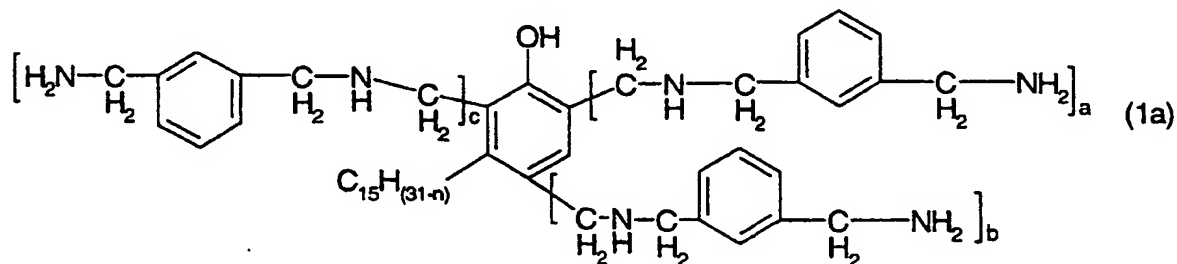


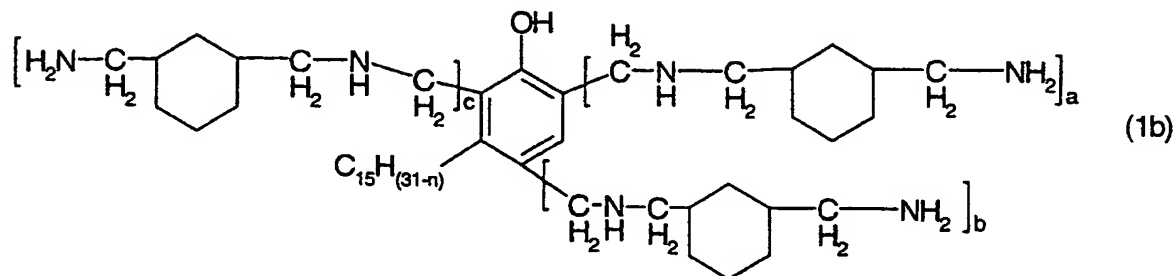
where n is 0, 2, 4 or 6.

14. A Mannich base reaction product according to claim 13 wherein the extract cashew nutshell liquid contains cardanol (I) and of cardol (II) in a weight ratio of about 90:10 to about 98:2.

15. A Mannich base reaction product according to claim 13 obtained by combining the extract and the at least one aldehyde compound with a polyamine selected from xylenediamine, 1,3-bis(aminomethyl)cyclohexane, and mixtures thereof.

16. A Mannich base reaction product according to claim 15 comprising at least one compound represented by structural formulae (1a) or (1b):





wherein n has been defined above, a, b and c, are, independently of one another, 1 or 0, and an average value for the sum of (a+b+c) is about 1.2 for all of the compounds corresponding to formulae (1a) and/or (1b) in said Mannich base reaction product mixture.

17. An epoxy resin composition comprising:

- a) a compound according to claim 1 and
- b) an epoxy resin having on average more than one glycidyl group per molecule.

18. An epoxy resin composition comprising:

- a) a Mannich base reaction product according to claim 16 and
- b) an epoxy resin having on average more than one glycidyl group per molecule.

19. A curable formulation comprising:

- a) an epoxy resin composition according to claim 17 and
- b) a pigment.

20. A curable formulation comprising:

- a) an epoxy resin composition according to claim 18 and
- b) a pigment.

21. A method for coating a surface having reduced yellowness index using a phenalkamine-containing composition that comprises applying a curable formulation according to claim 19 to the surface.

22. A method for coating a surface having reduced yellowness index using a phenalkamine-containing composition that comprises applying a curable formulation according to claim 20 to the surface.

INTERNATIONAL SEARCH REPORT

International Application No

PCT/EP 99/04323

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 C07C215/50 C08G59/40 C08G59/62

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C07C C08G

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	GB 2 152 925 A (UNIV BRUNEL) 14 August 1985 (1985-08-14) page 1, line 1-26	1,2,4,7
Y	page 2, line 31-40; claims 1-9; examples 1-6	1-22
Y	EP 0 042 617 A (HOECHST AG) 30 December 1981 (1981-12-30) the whole document	1-22
Y	DE 20 25 159 A (REICHOLD ALBERT CHEMIE AG) 23 December 1970 (1970-12-23) the whole document	1-20
Y	US 4 444 634 A (KEMPTER FRITZ E ET AL) 24 April 1984 (1984-04-24) column 4, line 60 -column 5, line 5	1-20
	--- -/--	

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents :

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"8" document member of the same patent family

Date of the actual completion of the international search

14 October 1999

Date of mailing of the international search report

20/10/1999

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

Rufet, J

INTERNATIONAL SEARCH REPORT

International Application No
PCT/EP 99/04323

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	EP 0 002 499 A (BASF AG) 27 June 1979 (1979-06-27) page 2, paragraph 4; claims 1-10 -----	1-20

Form PCT/ISA/210 (continuation of second sheet) (July 1992)

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/EP 99/04323

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
GB 2152925 A	14-08-1985	US 4697038 A	29-09-1987
EP 0042617 A	30-12-1981	DE 3023464 A	04-02-1982
		AT 12510 T	15-04-1985
		DE 3124232 A	17-02-1983
		JP 1183757 C	27-12-1983
		JP 57044621 A	13-03-1982
		JP 58017776 B	09-04-1983
		US 4399268 A	16-08-1983
DE 2025159 A	23-12-1970	AT 301184 B	15-07-1972
		BE 748631 A	16-09-1970
		CH 522016 A	30-04-1972
		FR 2052315 A	09-04-1971
		GB 1309454 A	14-03-1973
		JP 48025759 B	31-07-1973
		NL 7004848 A	22-12-1970
		SE 357568 B	02-07-1973
US 4444634 A	24-04-1984	DE 3123968 A	13-01-1983
		AT 19640 T	15-05-1986
		BR 8203530 A	07-06-1983
		EP 0068243 A	05-01-1983
		ES 513253 A	16-03-1983
		JP 58045217 A	16-03-1983
EP 0002499 A	27-06-1979	DE 2755908 A	21-06-1979
		US 4188312 A	12-02-1980